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NEW COMBINATION OF R,R-FORMOTEROL AND BUDESONIDE IN A PHARMACEUTICAL COMPOSITION
USEFUL FOR TREATING RESPIRATORY DISORDERS, SUCH AS ASTHMA, RHINITIS AND COPD

Field of the invention

5 This invention relates to improvement in the treatment of mild, moderate and severe
asthma and other respiratory disorders such as rhinitis and chronic obstructive pulmonary
disease (COPD). More particularly, it relates to the use of the steroidal anti-inflammatory
drug budesonide in combination with the strongly active R,R-enantiomer (preferably as the
fumarate dihydrate salt) of the long-acting bronchodilator formoterol (R,R;S,S) for the
10 treatment of respiratory disorders such as mild, moderate and severe asthma, rhinitis and
COPD, and to pharmaceutical compositions containing the two active ingredients.

Background of the invention

15 The recognition more than 10 years ago of the fundamentally inflammatory nature of
asthma led to the suggestions that control of the underlying airway inflammation could
provide the key to the control of asthma at all levels of severity. Nevertheless many
patients with asthma of most levels of severity still receive no regular anti-inflammatory
treatment and are treated only with intermittent or regular bronchodilator therapy.

20 Prophylactic therapy is typically provided by steroids such as beclomethasone dipropionate
(BDP), flunisolide, triamcinolone acetonide, dexamethasone, mometasone furoate,
fluticasone propionate and budesonide or by way of sodium cromoglycate or nedocromil
sodium.

25 Long-acting β_2 -agonists such as formoterol and salmeterol, have different properties from
short-acting ones such as terbutaline and salbutamol. These long-acting bronchodilators
have been regarded as add-on treatment to steroid therapy. However, the long-acting
agonists are considered an alternative to a further increase in the dosage of inhaled steroids.
The side-effects of the steroids could therefore be minimized. Therapy should be aimed at
30 controlling symptoms so that normal life is possible and at the same time provide basis for

treating the underlying inflammation. An interesting approach for this treatment strategy would be to combine a β 2-agonist with fast onset of action for symptom control together with an anti-inflammatory agent like a glucocorticosteroid.

5 The most common cause for poor control of asthma is poor compliance in the long-time management of chronic asthma, particularly with prophylactic treatment such as inhaled steroids, which do not give immediate symptom relief. Patients will readily take β 2-agonist inhalers, since these provide rapid onset of symptoms, but often do not take the prophylactic therapy, such as inhaled steroids, regularly because there is no immediate 10 symptomatic benefit.

Drug stereoisomerism is increasingly being recognized as an issue having clinical, research and regulatory implications. Differences in the pharmaco-dynamic and pharmacokinetic properties of stereoisomers are well documented e.g. the pharmacological properties of 15 drug enantiomers can be dramatically different; one isomer may be predominantly responsible for the desired therapeutic action and the other for the side effects. In the case of formoterol (a mixture of R,R and S,S), the R,R-enantiomer is about 1000 times more potent than the S,S-isomer (see Trofast et al (1991)).

20 Earlier mentioned combinations of long-acting β -agonists and steroids include the use of salmeterol/bclomethasone dipropionate (US 5,208,226, Glaxo), salmeterol/fluticasone propionate (US 5,270,305, Glaxo) and formoterol/budesonide (US 5,674,860, Astra). The inhaled route of administration enables the dose to be delivered directly to the airways. By this type of administration, it is possible to give a small dose and thereby minimizing 25 unwanted side-effects.

Summary of the invention

It has now surprisingly been found that a combination of R,R formoterol and budesonide can be used for the treatment of respiratory disorders such as asthma, rhinitis and COPD.

5 According to the invention there is provided a pharmaceutical combination which comprises R,R formoterol in combination with budesonide.

Detailed description of the invention

10 The present invention provides a novel combination therapy using the long-acting bronchodilator R,R-formoterol (preferably as the fumarate dihydrate salt) and the glucocorticosteroid budesonide.

In a first aspect the present invention provides a pharmaceutical combination which 15 comprises:

- (a) R,R-formoterol, or a pharmaceutical acceptable salt or solvate thereof,
- (b) budesonide; and optionally
- (c) one or more pharmaceutically acceptable additives, diluents or carriers;

20 Preferably the molar ratio of (a) to (b) is from 1:4 to 1:100.

The word "combination" is used to describe the invention because the components can be administered simultaneously or sequentially for use in therapy. Thus the active ingredients 25 (a) and (b) are not necessarily, but may be, used as an admixture, they still have the desired effect if they are administered sequentially or separately. Preferably they are not administered more than about two hours apart, for example no more than 30 minutes apart.

The first main ingredient of the combination of the invention is the single enantiomer R,R-formoterol i.e. R,R-(N-[2-hydroxy-5-[1-hydroxy-2-[[2-(4-methoxyphenyl)-1-methyl-30 ethyl]-amino]-ethyl]phenyl]-formamide, an adrenoceptor agonist which selectively

stimulates β_2 -receptors, thus producing relaxation of bronchial smooth muscle, inhibition of the release of endogenous spasmogens, inhibition of oedema caused by endogenous mediators, and increased mucociliary clearance. The compound can be prepared by methods described in "Large-Scale Synthesis of Enantio- and Diastereomerically Pure

5 (R,R)-formoterol" by R. Hett et al. in *Organic Process Research & Development*, 2 (1998), 96-99 or in "Steric Aspects of Agonism and Antagonism at β -adrenoceptors: Synthesis of and Pharmacological Experiments With the Enantiomers of Formoterol and Their Diastereomers" by J. Trofast et al in *Chirality* 3 (1991), 443-450.

10 The other main ingredient is budesonide i.e. 16,17-butyridenebis(oxy)-11,21-dihydroxy-pregna-1,4-diene-3,20-dione. The compound can be prepared by the methods described in US 3,929,768. The compound exists as epimers, and either epimer can be used in the combinations of the invention, including the 22R epimer.

15 A combination, preferably a fixed combination i.e. given in admixture, of the compounds of the invention will establish a higher compliance for patients and it provides a rescue medicine thereby avoiding the necessity for the patient of carrying two different inhalers. This simplifies the life for the patients considerably and makes life more comfortable and secure.

20

According to another aspect of the invention there are provided pharmaceutical compositions comprising effective amounts of R,R-formoterol (and/or physiologically acceptable salt and/or solvate thereof) and budesonide as a preparation for simultaneous, sequential or separate administration by inhalation in the treatment of respiratory disorders 25 such as asthma, rhinitis and COPD. Reference to formoterol and salts and solvates thereof includes all combinations of solvates and salts of formoterol such as solvates of salts.

The invention additionally relates to the use of R,R-formoterol (and/or a physiologically acceptable salt and/or solvate thereof) and budesonide in the manufacture of 30 pharmaceutical compositions as preparations for simultaneous, sequential or separate

administration of R,R-formoterol and budesonide by inhalation in the treatment of respiratory disorders such as asthma, rhinitis and COPD.

According to a further feature of the invention there is provided a method of treating respiratory disorders which comprises the simultaneous, sequential or separate administration by inhalation of effective amounts of R,R-formoterol (and/or a physiologically acceptable salt and/or solvate thereof) and budesonide.

Suitable physiological salts of R,R-formoterol include acid addition salts derived from inorganic and organic acids, such salts as the chloride, bromide, sulphate, phosphate, maleate, fumarate, tartrate, citrate, benzoate, 4-methoxybenzoate, 2- or 4-hydroxybenzoate, 4-chlorobenzoate, p-toluene-sulphonate, methanesulphonate, ascorbate, salicylate, acetate, succinate, lactate, glutarate, gluconate, tricarballylate, hydroxynaphthalene carboxylate or oleate. R,R-Formoterol is preferably used in the form of its fumarate salt and as a dihydrate of that salt.

The intended dose regimen is once or twice a day, where the suitable daily dose of R,R-formoterol is in the range of from about 5 to about 250 nmol (preferably from about 10 to about 120 nmol) and for budesonide a daily dose of about 0.1 μ mol to about 3 μ mol with a preferred dose of about 0.1 μ mol to about 2 μ mol. The doses of R,R-formoterol to budesonide should be selected to be within the molar range of from 1:4 to 1:100. The two drugs may be administered separately in the same ratio. The dose of choice will strongly depend on the patient (age, weight etc) and the severity of the disease (mild, moderate, severe asthma etc).

25

The combination is inhaled from a nebulizer, from a pressurized metered dose inhaler or as a dry powder from a dry powder inhaler e.g. multidose reservoir systems from Astra (Turbuhaler[®]) or from a dry powder inhaler utilizing gelatine, plastic or other capsules, cartridges or blister packs. A diluent or carrier, generally being non-toxic and chemically inert to the medicament e.g. lactose, dextran, mannitol or glucose or any additives that will

give the medicament a certain taste can be added to the powdered medicament in an amount of from 50 µg to 25 mg per dose, more preferably in an amount of from 50 µg to 10 mg, most preferably in an amount of from 100 to 2000 µg.

- 5 One or more of the ingredients is preferably in the form of a dry powder, more preferably a micronized dry powder, most preferably an agglomerated micronized dry powder. As an alternative to agglomeration, the finely divided active ingredients may be in the form of an ordered mixture with the pharmaceutically acceptable additive, diluent or carrier. An ordered mixture comprises fine particles of an active ingredient in association with coarse
- 10 particles of the pharmaceutically acceptable additive, diluent or carrier. A fraction of fine particles of carrier may also be present. The ingredients used in the invention can be obtained in these preferred forms using methods known to those skilled in the art. The particle size of the active ingredients is less than 20 µm, preferably less than 10 µm.
- 15 When the ingredients of the system are adapted to be administered from a pressurized inhaler, they are preferably in micronized form. They are dissolved, or, preferably suspended in a liquid propellant mixture. The propellants which can be used include chlorofluorocarbons, hydrocarbons or hydrofluoroalkanes. Especially preferred propellants are P134a (tetrafluoroethane) and P227 (heptafluoropropane) each of which may be used
- 20 alone or in combination. They are optionally used in combination with one or more other propellants and/or one or more surfactants and/or one or more other excipients, for example ethanol, a lubricant, an anti-oxidant and/or a stabilising agent.

When the ingredients of the system of the invention are adapted to be administered via a

- 25 nebuliser they may be in the form of a nebulised aqueous suspension or solution, with or without a suitable pH or tonicity adjustment, either as a unit dose or multidose device.

The invention is illustrated by the following examples which are not intended to limit the scope of the application. In the examples micronization is carried out such that the particle size range for each component is suitable for administration by inhalation. The dry powder

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formulation containinig an additive, diluent or carrier could be either in agglomerated form or as ordered mixtures .

Example 1.

Per dose

	R,R-Formoterol fumarate dihydrate	6 µg
5	Budesonide	100 µg

Example 2.

	R,R-Formoterol fumarate dihydrate	6 µg
10	Budesonide	200 µg

Example 3.

	R,R-Formoterol fumarate dihydrate	3 µg
15	Budesonide	100 µg

Example 4.

	R,R-Formoterol fumarate dihydrate	3 µg
20	Budesonide	50 µg
	Lactose monohydrate	up to 0.5, 1,5,10,20 mg

Example 5.

25	R,R-Formoterol fumarate dihydrate	3 µg
	Budesonide	100 µg
	Lactose monohydrate	up to 0.5, 1, 5, 10, 20 mg

Example 6.

R,R-Formoterol fumarate dihydrate	3 µg
Budesonide	200 µg
Lactose monohydrate	up to 0.5, 1, 5, 10, 20 mg

5 **Example 7.**

R,R-Formoterol fumarate dihydrate	3 µg
Budesonide	100 µg
Oleic acid (based on propellant)	0.005 %
10 Ethanol (based on propellant)	1.5 %
Propellant P134a	up to 25, 50 or 100 µl

Example 8.

R,R-Formoterol fumarate dihydrate	6 µg
15 Budesonide	200 µg
Oleic acid (based on propellant)	0.01 %
Ethanol (based on propellant)	1.5 %
Propellant P227/P134a (15/85)	up to 25, 50 or 100 µl

20 **Example 9.**

2.6 parts of R,R-formoterol fumarate dihydrate and 896.8 parts of lactose monohydrate were mixed in a tumbling mixer to an evenly distributed mixture, whereafter the mixture was micronized in a spiral jet mill using a pressure and feeding suitable to obtain a particle size of less than 3 um. The micronized particles were then treated using a method described in WO 95/05805 to remove amorphous regions in their crystal structure. 98 parts of micronized budesonide were added and the mixture was remicronized at a lower pressure in a spiral jet mill to a homogeneous mixture. The powder was then agglomerated by feeding into a screw feeder (K-tron), sieved, spheronized in a rotating pan, then sieved

again, spheronized once more before final sieving (0.8 mm mesh size) to give a powder suitable for an inhaler.

Example 10.

5 Example 9 was repeated with identical conditions but using 2.6 parts of micronized R,R-formoterol fumarate dihydrate, 798.8 parts of micronized lactose monohydrate and 196 parts of micronized budesonide.

Claims.

1. A pharmaceutical combination which comprises:
 - 5 (a) R,R-formoterol, or a pharmaceutical acceptable salt or solvate thereof,
 - (b) budesonide; and optionallyone or more pharmaceutically acceptable additives, diluents or carriers.
- 10 2. A pharmaceutical combination according to claim 1 wherein the molar ratio of (a) to (b) is from 1:4 to 1:100.
- 15 3. A pharmaceutical combination according to claim 1 or 2 in which the R,R-formoterol is in the form of the fumarate dihydrate salt.
4. A pharmaceutical combination according to any one of claims 1 to 3 in which the combination is fixed and given in admixture.
- 20 5. A pharmaceutical combination according to any one of claims 1 to 4 in a form suitable for administration from a pressurised inhaler.
- 25 6. A pharmaceutical combination according to claim 5 comprising R,R-formoterol, or a pharmaceutical acceptable salt or solvate thereof, budesonide; and optionally a propellant and one or more other surfactants and/or one or more excipients.
7. A pharmaceutical combination according to claim 6 in which the propellant is HFA 227.
- 30 8. A pharmaceutical combination according to any one of claims 1 to 7 for use for the treatment or prophylaxis of a respiratory disorder.

9. A pharmaceutical combination according to any one of claims 1 to 7 for use for the treatment or prophylaxis of asthma, rhinitis or COPD.

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INTERNATIONAL SEARCH REPORTInternational application No.
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A. CLASSIFICATION OF SUBJECT MATTER

IPC7: A61K 31/58, A61K 31/165

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 9815280 A1 (ASTRA AKTIEBOLAG ET AL), 16 April 1998 (16.04.98) --	1-19
A	CHIRALITY, Volume 3, 1991, Trofast, Jan et al, "Steric Aspects of Agonism and Antagonism at Beta-Adrenoceptors:" page 443 - page 450 -- -----	1-19

 Further documents are listed in the continuation of Box C. See patent family annex.

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Patent document cited in search report	Publication date	Patent family member(s)		Publication date
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